

Substitute Form PTO-1449 Modified	U.S. Department of Commerce Patent and Trademark Office	Attorney's Docket No. 07039-784US1	Application No. 10/568,414
Information Disclosure Statement by Applicant (Use several sheets if necessary) (37 CFR §1.808)		Applicant Jan O. Aasly et al.	
		Filing Date December 7, 2006	Group Art Unit 1632

U.S. Patent Documents

Examiner Initial	Desig. ID	Document Number	Publication Date	Patentee	Class	Subclass	Filing Date If Appropriate

Foreign Patent Documents or Published Foreign Patent Applications

Examiner Initial	Desig. ID	Document Number	Publication Date	Country or Patent Office	Class	Subclass	Translation	
							Yes	No
/JH/	1.	WO 06/68492	06/29/06	WIPO				

Other Documents (include Author, Title, Date, and Place of Publication)

Examiner Initial	Desig. ID	Document
/JH/	2.	GenBank Accession Number AY792511 dated 11/15/04, 5 pages
	3.	Bonifati et al., "Mutations in the <i>DJ-1</i> Gene Associated with Autosomal Recessive Early-Onset Parkinsonism," <u>Science</u> , 2003, 299:256-259
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	5.	Chartier-Harlin et al., "α-synuclein locus duplication as a cause of familial Parkinson's disease," <u>Lancet</u> , 2004, 364:1167-1169
	6.	Davies et al., "Mutations of the <i>BRAF</i> gene in human cancer," <u>Nature</u> , 2002, 417:949-954
	7.	de Rijk et al., "Prevalence of Parkinson's disease in the elderly: the Rotterdam Study," <u>Neurology</u> , 1995, 45:2143-2146
	8.	Dibb et al., "Switching on kinases: oncogenic activation of BRAF and the PDGFR family," <u>Nat. Rev. Cancer</u> , 2004, 4:718-727
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	11.	Funayama et al., "A New Locus for Parkinson's Disease (<i>PARK8</i>) Maps to Chromosome 12p11.2-q13.1," <u>Ann. Neurol.</u> , 2002, 51:296-301
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	15.	Kitada et al., "Mutations in the <i>parkin</i> gene cause autosomal recessive juvenile parkinsonism," <u>Nature</u> , 1998, 392:605-608
	16.	Kong and Cox, "Allele-Sharing Models: LOD Scores and Accurate Linkage Tests," <u>Am. J. Hum. Genet.</u> , 1997, 61:1179-1188
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Examiner Signature	Date Considered
EXAMINER: Initials citation considered. Draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant.	

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(37 CFR §1.98(b))			

Other Documents (include Author, Title, Date, and Place of Publication)		
Examiner Initial	Desig. ID	Document
/JH/	18.	Lander and Kruglyak, "Genetic dissection of complex traits: guidelines for interpreting and reporting linkage results," <u>Nat. Genet.</u> , 1995, 11:241-247
↓	19.	Lang and Lozano, "Parkinson's Disease. First of Two Parts," <u>New Engl. J. Med.</u> , 1998, 339:1044-1053
	20.	Mata et al., "Parkin genetics: one model for Parkinson's disease," <u>Hum. Mol. Genet.</u> , 2004, 13:R127-R133
	21.	Paisán-Ruiz et al., "Cloning of the Gene Containing Mutations that Cause <i>PARK8</i> -Linked Parkinson's Disease," <u>Neuron</u> , 2004, 44:595-600
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	27.	Spillantini et al., "α-Synuclein in Lewy bodies," <u>Nature</u> , 1997, 388:839-840
	28.	Tanner et al., "Parkinson Disease in Twins: An Etiologic Study," <u>JAMA</u> , 1999, 281(4):341-346
	29.	Valente et al., "Hereditary Early-Onset Parkinson's Disease Caused by Mutations in <i>PINK1</i> ," <u>Science</u> , 2004, 304:1158-1160
	30.	Vila and Przedborski, "Genetic clues to the pathogenesis of Parkinson's disease," <u>Nat. Med.</u> , 2004, 10 Suppl:S58-S62
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	32.	Zarranz et al., "The New Mutation, E46K, of α-Synuclein Causes Parkinson and Lewy Body Dementia," <u>Ann. Neurol.</u> , 2004, 55:164-173
	33.	Zimprich et al., "Mutations in <i>LRRK2</i> Cause Autosomal-Dominant Parkinsonism with Pleomorphic Pathology," <u>Neuron</u> , 2004, 44:601-607
	34.	Zimprich et al., "The <i>PARK8</i> locus in autosomal dominant parkinsonism: confirmation of linkage and further delineation of the disease-containing interval," <u>Am. J. Hum. Genet.</u> , 2004, 74:11-19

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